# **AMENDMENTS TO THE CLAIMS**

# **Listing of Claims**

This listing of the claims will replace all prior versions, and listings, of claims in this application.

#### 1-59. (Cancelled)

- 60. (Currently Amended) A method of identifying compounds that bind to a leukotriene  $A_4$  (LTA<sub>4</sub>) hydrolase comprising the amino acid sequence of SEQ ID NO:1, the method comprising the steps of:
- (a) crystallizing a purified LTA<sub>4</sub> hydrolase <del>comprising the amino acid sequence of SEQ ID NO:1 together with bestatin to form an LTA<sub>4</sub> hydrolase crystal, wherein crystallization is performed as liquid liquid diffusion in a capillary using equal volumes of a buffer: enzyme solution comprising:</del>

i) a buffer solution comprising about 28% PEG8000, about 0.1 M Na-acetate, about 0.1 M imidazole at a pH of about 6.8 and with about 5 mM YbCl<sub>3</sub> as an additive; and

ii) an enzyme solution comprising about 5 mg/ml LTA<sub>4</sub> hydrolase comprising the amino acid sequence of SEQ ID NO:1 in about 10 mM Tris-HCl at a pH of about 8, supplemented with about 1 mM bestatin;

- (b) determining the atomic coordinates of said LTA<sub>4</sub> hydrolase crystal; and
- (c) screening the atomic coordinates of a set of candidate compounds against the atomic coordinates of said LTA<sub>4</sub> hydrolase crystal obtained in step a) to identify compounds that bind to the LTA<sub>4</sub> hydrolase.
- 61. (**Previously Presented**) The method of claim 60, wherein the LTA<sub>4</sub> hydrolase is purified by adsorption chromatography on hydroxyapatite and anion-exchange chromatography.

#### 62-67. (**Cancelled**)

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68. (**Previously Presented**) The method of claim 60, wherein the atomic coordinates of said LTA<sub>4</sub> hydrolase crystal correspond to the atomic coordinates defining atom 1 to atom 4876 as set forth in Table 9.

#### 69. (Cancelled)

- 70. (**Currently Amended**) A method of designing an inhibitor or agonist of LTA<sub>4</sub> hydrolase comprising the amino acid sequence of SEQ ID NO:1, the method comprising the steps of:
- (a) crystallizing a purified LTA<sub>4</sub> hydrolase comprising the amino acid sequence of SEQ ID NO:1 together with bestatin to form a crystal and thereafter determining its conformational structure, wherein crystallization is performed as liquid liquid diffusion in a capillary using equal volumes of a buffer: enzyme solution comprising:
- <u>i)</u> a buffer solution comprising about 28% PEG8000, about 0.1 M Na-acetate, about 0.1 M imidazole at a pH of about 6.8 and with about 5 mM YbCl<sub>3</sub> as an additive; and
- ii) an enzyme solution comprising about 5 mg/ml LTA<sub>4</sub> hydrolase comprising the amino acid sequence of SEQ ID NO:1 in about 10 mM Tris-HCl at a pH of about 8, supplemented with about 1 mM bestatin;
- (b) identifying at least one compound that is at least in part complementary to the LTA<sub>4</sub> hydrolase by the use of the conformational structure of the crystal complex obtained in step a);
- (c) soaking the crystallized LTA<sub>4</sub> hydrolase obtained in step a) with a solution of a compound identified in step b) to obtain a complex of the crystal of said LTA<sub>4</sub> hydrolase and said compound; and
- (d) performing X-ray crystallography of the crystal complex of LTA<sub>4</sub> hydrolase and said compound to determine the structure thereof, thereby identifying the compound as an inhibitor or agonist of LTA<sub>4</sub> hydrolase.
- 71. (**Previously Presented**) The method of claim 70, wherein the LTA<sub>4</sub> hydrolase is purified by adsorption chromatography on hydroxyapatite and anion-exchange chromatography.

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72. (**Previously Presented**) The method of claim 70, wherein said compound is an inhibitor of LTA<sub>4</sub> hydrolase.

### 73-75. (**Cancelled**)

76. (**Previously Presented**) The method of claim 70, wherein the atomic coordinates of said LTA<sub>4</sub> hydrolase crystal correspond to the atomic coordinates defining atom 1 to atom 4876 as set for in Table 9.

# 77. (Cancelled)

- 78. (**Previously Presented**) The method of claim 70, further comprising the step of refining the structure of said compound obtained in step d) via computer modeling using data obtained from the X-ray crystallography in step d) and repeating steps b)-d).
- 79. (**Previously Presented**) The method of claim 70, wherein the complex obtained in step c) comprises bestatin.

80-86. (Cancelled)